This listing of claims presented below replaces all prior versions and listings of claims in the application.

Listing of Claims

IN THE CLAIMS

- 1. (Currently Amended) A method of diagnosis diagnosing of immunological recurrent spontaneous abortion, characterized by in vitro comprising the steps of: determining the <u>a</u> level of antinuclear antibody in a body fluid sample from the <u>a</u> patient <u>by using an isolated human chromosome No. 2 as antigen; and comparing the result with the determined level with a level of the corresponding antinuclear antibody of normal <u>a</u> control.</u>
- 2. (Cancelled)
- 3. (Cancelled)
- 4. (Cancelled)
- 5. (Cancelled)
- 6. (Cancelled)
- 7. (Currently Amended) A kit for the diagnosis of diagnosing immunological recurrent spontaneous abortion, comprising an isolated human chromosome No. 2 or fragment thereof containing fibronectin encoding gene derived from male (s) as antigen.
- 8. (Cancelled)
- 9. (Cancelled)
- 10. (Cancelled)
- 11. (Cancelled)
- 12. (Cancelled)
- 13. (Original) The kit according to claim 7, wherein said antigen is coated on a solid carrier.

- 14. (Currently Amended) The kit according to claim13, further comprising an enzyme-labeled secondary antibody, necessary suitable buffer solutions, and instructions.
- 15. (Currently Amended) A method for monitoring the <u>a</u> therapeutic effect for immunological recurrent spontaneous abortion, characterized by in vitro comprising the steps of:

determining the <u>a</u> level of <u>an</u> antinuclear antibody in a body fluid sample of the <u>a</u> patient, after <u>a</u> treatment, <u>by using an isolated human chromosome No. 2 as antigen;</u> and comparing the <u>result determined level</u> with the corresponding level before <u>the</u> treatment.

- 16. (Cancelled)
- 17. (Cancelled)
- 18. (Cancelled)